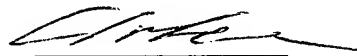


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For and on behalf of RWS Group Ltd

The 2nd day of April 2009

FEDERAL REPUBLIC OF GERMANY



**Priority Certificate
DE 102 51 729.0
for the filing of a Patent Application**

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Applicant/Proprietor: BASF Coatings AG, 48165 Münster/DE

Title: UV-active binders

IPC: C 08 F 220/10, C 08 F 2/46, C 09 D 133/10

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UV-active binders

The invention relates to a process for preparing
5 radiation-curable and/or dual-cure poly(meth)acrylates,
to the poly(meth)acrylates themselves, to their use as
a component in the preparation of dispersions or as a
component in coating formulations, to coating
formulations comprising the poly(meth)acrylates of the
10 invention and to a process for preparing the coating
formulations and their use.

UV-curable and dual-cure poly(meth)acrylates are of
particular interest for use in topcoats. Poly-
15 (meth)acrylates generally have excellent outdoor
weather stabilities. In conjunction with a UV curing
technology, additional advantages are achievable. For
example, the scratch resistance of the coating can be
increased significantly, with an attendant improvement
20 in coating performance. Particularly important,
however, are improvements in application, in particular
a very rapid drying of the coating materials. This
property is critical for rapid processing technologies.

25 In accordance with the prior art, UV-curable
polyacrylates are reacted by copolymerizing glycidyl
methacrylate followed by thermal reaction with acrylic
acid in the presence of a catalyst (DE-A 2 436 186,
EP-A 0 650 978). Disadvantages associated with this
30 prior art mode of preparation are the attendant
secondary reactions and deteriorations in color. In

view of the reaction conditions, particularly the high temperatures, under which the reaction is carried out it is absolutely necessary to use stabilizers in order to prevent the free-radical polymerization of the acrylic acid used.

A conventional acidic esterification of poly(meth)acrylates containing hydroxy-functional side chains with acrylic acid is not possible, since in the course of such reaction the ester bonds of the poly(meth)acrylate are cleaved.

Known from the prior art is the functionalization of polymeric compounds with (meth)acrylic acid and/or (meth)acrylic esters.

EP-A 0 999 230 and EP-A 0 999 229 relate to processes for preparing (meth)acrylic esters of hydroxy-functional siloxanes and/or polyalkylene-modified siloxanes ('230) and also of polyoxyalkylenes ('229) by esterification or transesterification of the siloxanes or polyoxyalkylenes, respectively, with (meth)acrylic acid and/or (meth)acrylic esters in the presence of an enzyme. According to EP-A 0 999 230 and EP-A 0 999 229, however, only the specific polymers referred to are reacted with (meth)acrylic acid and/or (meth)acrylic esters. There is no mention of a reaction of poly(meth)acrylates.

E. Marechal et al., Polymer Bulletin 26, 55 to 62 (1991) relates to the transesterification of oligo(methacrylates) containing terminal ester groups with allyl alcohol in the presence of lipase. Transesterification takes place only at the terminal groups.

H. Ritter et al., Polymer Bulletin 21, 535 to 540 (1989) relates to the lipase catalyzed acetylation of

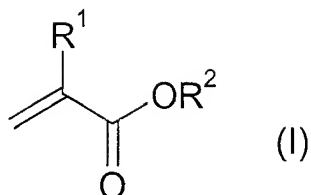
methacrylic acid polymers containing OH groups. The acetylation takes place in the presence of vinyl acetate. Reaction with vinyl acetate produces a very good leaving group, with the aldehyde formed being
5 easily removable from the reaction mixture. Nevertheless, the reaction time amounts to 2 to 15 days.

H. Ritter et al., Makromol. Chem. 193, 323 to 328
10 (1992) relates to the enzymatically catalyzed acylation of OH-containing comblike methacrylic acid polymers with active esters, such as vinyl acetate, phenyl acetate, 4-fluorophenyl acetate, and phenyl stearate. There is no mention of an esterification of the
15 polymers with acrylates. The reaction times of the reaction according to Ritter et al. are very long (2, 4 and 6 days).

The object of the present invention is to provide a
20 gentle and selective process for preparing poly(meth)acrylates functionalized with (meth)acrylic acid and/or (meth)acrylates which is more variable than the known preparation process starting from glycidyl methacrylate, is able to start from less expensive
25 starting substances, and allows a gentler preparation, so that new kinds of poly(meth)acrylates substituted by (meth)acrylic groups are obtainable.

The object is achieved by a process for preparing UV-
30 curable and/or dual-cure poly(meth)acrylates, comprising the following steps:

- a) preparing a poly(meth)acrylate containing hydroxy-functional side chains by polymerizing
35 aa) at least one (meth)acrylate of the general formula (I) as component A

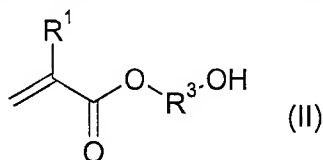


in which

R^1 is H, CH_3 or CH_2OH and

R^2 is an alkyl or cycloalkyl radical which is unsubstituted or substituted by functional groups such as acrylic, ether, amino, epoxy, halogen or sulfonic acid groups, preferably a C_1 to C_{18} alkyl radical, more preferably a C_1 to C_8 alkyl radical, very preferably a C_1 to C_8 alkyl radical unsubstituted by functional groups, in particular a methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, 2-ethylhexyl, tert-butyl, cyclohexyl, tert-butylcyclohexyl, isobornyl or trimethylcyclohexyl radical; and

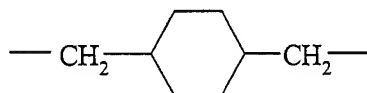
ab) at least one hydroxyalkyl (meth)acrylate of the general formula (II) as component B



in which

R^1 is H, CH_3 or CH_2OH and

R^3 is $-(\text{CH}_2)_n-$, $-\text{CH}_2-\text{CH}(\text{CH}_3)-\text{CH}_2-$ or $-\text{CH}_2\text{CH}(\text{CH}_3)-$ or $-\text{CH}(\text{CH}_3)\text{CH}_2-$ or



n is at least 2, preferably 2 to 8, more preferably 2 to 6, very preferably 2 to 4,

the hydroxyalkyl (meth)acrylate of the
general formula (II) being selected in
particular from the group consisting of
5 2-hydroxyethyl (meth)acrylate,
2-hydroxypropyl (meth)acrylate or
hydroxybutyl (meth)acrylate; and

ac) if desired, further comonomers,
10 copolymerizable with the (meth)acrylates of
the general formula (I) and (II), as
component C, preferably selected from the
group consisting of styrene, acrylonitrile,
vinyl acetate, vinyl propionate, vinyl
15 chloride, vinylidene chloride, butadiene, and
adducts of Versatic acid glycidic residues and
unsaturated acids, especially (meth)acrylic
acid, and

20 ad) if desired, auxiliary monomers as component D
preferably selected from the group consisting
of (meth)acrylic acid, itaconic acid, maleic
acid, fumaric acid, crotonic acid, and the
amides of said acids;

25

and

b) transesterifying or esterifying the poly-
(meth)acrylate containing hydroxy-functional side
30 chains with a (meth)acrylate or (meth)acrylic
acid, preferably with methyl, ethyl, 2-ethylhexyl
or butyl (meth)acrylate, in the presence if
desired of stabilizers selected from the group
consisting of 2,6-dibutylphenols such as di-tert-
35 butylphenol, p-cresol, hydroquinone, dimethyl-
hydroquinone, phenothiazines and phosphorous
esters, in the presence of an enzyme which

catalyzes the transesterification or esterification.

"(Meth)acrylic acid" is used as an abbreviation for
5 "methacrylic acid or acrylic acid"; correspondingly,
"(meth)acrylate" is used as an abbreviation for
"methacrylate or acrylate".

The process of the invention enables functional groups,
especially (meth)acrylic groups, to be introduced
10 gently into poly(meth)acrylates without the likelihood
of cleavage of the ester groups of the poly-
(meth)acrylate. Moreover, it is possible to prepare the
functionalized polymers starting from poly(meth)-
acrylates containing hydroxy-functional side chains,
15 which are substantially less expensive than glycidyl-
functionalized poly(meth)acrylates used hitherto in the
prior art.

Step a)

20

In a preferred embodiment of the process of the
invention step a) is carried out using

- 10 to 80% by weight, preferably 20 to 80% by
weight, more preferably 30 to 70% by weight of
25 component A, and
- 10 to 80% by weight, preferably 20 to 70% by
weight, more preferably 20 to 60% by weight of
component B, and
- 0 to 50% by weight, preferably 0 to 40% by weight,
30 more preferably 5 to 25% by weight of component C,
and
- 0 to 15% by weight, preferably 0 to 10% by weight,
more preferably 0.5 to 5% by weight of
component D.

35

The poly(meth)acrylates used in accordance with the
invention, containing hydroxy-functional side chains,
can be prepared by various methods known to the skilled

worker. Preference is given to their preparation by free-radical polymerization.

5 The polymerization takes place in general by emulsion, solution or bulk polymerization, preference being given to emulsion polymerization or solution polymerization.

10 In one embodiment the poly(meth)acrylates containing hydroxy-functional side chains are prepared by emulsion polymerization. In the case of emulsion polymerization components A, B, and, where appropriate, C and, where appropriate, D are reacted with one another in the presence of water, emulsifiers, initiators, and, where appropriate, regulators.

15

Emulsifiers used are generally anionic, nonionic, cationic or amphoteric emulsifiers, with anionic or nonionic emulsifiers being preferred. Suitable anionic emulsifiers are sodium, potassium or ammonium salts of
20 long-chain aliphatic carboxylic acids and sulfonic acids, alkali metal C₁₂₋₁₆ alkyl sulfates, oxethylated and sulfated or sulfonated long-chain aliphatic alcohols or alkylphenols, and sulfodicarboxylic esters. Suitable nonionic emulsifiers are oxethylated fatty
25 alcohols and alkylphenols, the ethylene oxide units possibly amounting to between 2 and 50 mol/mol. Suitable cationic emulsifiers are ammonium, phosphonium, and sulfonium compounds including at least one long aliphatic hydrocarbon chain as a hydrophobic
30 moiety. It is also possible to use a combination of different emulsifiers: for example, ionic and nonionic emulsifiers.

35 The water used is preferably distilled or deionized, since salts may affect the stability of the emulsion. Generally speaking, the polymerization process is carried out under nitrogen, since oxygen inhibits the polymerization.

The molecular weight of the poly(meth)acrylates containing hydroxy-functional side chains can be lowered by adding regulators. Suitable regulators are, 5 for example, halogenated compounds such as carbon tetrachloride, carbon tetrabromide, bromal, benzyl bromide, and trichlorobromomethane, or mercaptans such as butyl mercaptan or dodecyl mercaptan or Rongalit® C. Suitable initiators are in general all initiators known 10 to the skilled worker for polymerizing (meth)acrylates. Use is made in general of water-soluble peroxo compounds such as alkali metal or ammonium persulfate, hydrogen peroxide or tert-butyl peroxyethylhexanoate. Also suitable are redox systems such as H₂O₂-ascorbic 15 acid, H₂O₂-Fe(II)/Fe(III), H₂O₂-Ce(IV), persulfites-Fe, metabisulfites-Fe or hydroperoxides-metal salts. The initiators are used generally in an amount of 0.05 to 8% by weight, preferably 0.2 to 2% by weight, based on amount of monomers used.

20 Any initiators still present after the polymerization can be deactivated after the polymerization in order to prevent possible polymerization of the poly-(meth)acrylates prepared in accordance with the 25 invention in step b). The deactivation is generally accomplished by adding a reducing agent, e.g., ascorbic acid.

The polymerization is generally conducted within a 30 temperature range from 30 to 120°C, preferably 40 to 110°C, more preferably 50 to 90°C. The polymerization is generally conducted under a pressure of 1 to 20, preferably 1 to 15 bar, more preferably 1 to 5 bar.

35 The emulsifiers are generally used in an amount of 0.5 to 15% by weight, preferably of 0.5 to 10% by weight, more preferably 0.5 to 5% by weight, based on the

amount of components A, B, if desired, C, and, if desired, D that are used.

5 The particle diameter of the poly(meth)acrylates containing hydroxy-functional side chains that are obtained after polymerization is generally 20 to 1000 nm, preferably 20 to 500 nm, more preferably 50 to 400 nm, determined by means of light scattering.

10 The pH during the emulsion polymerization is generally between 1 and 6, preferably between 2 and 6. The hydroxyl numbers are generally at least 20 to 180, preferably at least 40 to 120. The solids content of the dispersions is generally 10 to 50, preferably 20 to
15 40, and the glass transition temperature of the polymers obtained is generally between -40 and +80°C.

The resultant poly(meth)acrylates containing hydroxy-functional side chains generally have an average
20 molecular weight of 1000 to 2 000 000, preferably 1000 to 1 000 000, more preferably 50 000 to 500 000. The average molecular weight was determined by means of gel permeation chromatography (GPC). The molecular weight in question is the number-average molecular weight.

25

The poly(meth)acrylates containing hydroxy-functional side chains can be prepared by means of a one-pot or batch procedure, feed techniques, and continuous procedures. The conduct of said procedures is known to
30 the skilled worker.

The poly(meth)acrylate containing hydroxy-functional side chains that is obtained in step a) can be isolated by methods known to the skilled worker. One embodiment
35 is described, for example, in EP-A 0 029 637, but in the process according to the present specification hydroxyl-free solvents are used, or in a second step a hydroxyl-containing solvent is replaced by a hydroxyl-

free solvent. For use in step b) of the process of the invention the poly(meth)acrylate containing hydroxy-functional side chains, following its isolation, is used in water-free form.

5

In another preferred embodiment the poly(meth)acrylates containing hydroxy-functional side chains are prepared by solution polymerization. In the solution polymerization the components A, B, and, where appropriate, C and, where appropriate, D are reacted with one another in the presence of a solvent, initiator and, where appropriate, regulators.

Initiators suitable for the solution polymerization are peroxides such as dialkyl peroxides, e.g., di-tert-butyl peroxide and di-tert-amyl peroxide, peroxy esters such as tert-butyl peroxy-2-ethylhexanoate and tert-amyl peroxy-2-ethylhexanoate, diacyl peroxides such as benzoyl peroxide, lauroyl peroxide, and decanoyl peroxide, percarbonates such as tert-butyl peroxyisopropyl carbonate, di-2-ethylhexyl peroxydicarbonate, perketals and ketone peroxides, and also azo initiators such as 2,2'-azobis(2,4-dimethylpentanenitrile), 2,2'-azobis(2-methylpropanonitrile), 2,2'-azobis(2-methylbutanonitrile), 1,1'-azobis(cyclohexanecarbonitrile), 2,2'-azobis(2,4,4-trimethylpentane), and 2-phenylazo-2,4-dimethyl-4-methoxyvaleronitrile.

Preferred solvents are those not disruptive to an enzymatic reaction in accordance with step b), so that removal of the solvent prior to execution of step b) is unnecessary. Particular preference is given to solvents selected from methyl isobutyl ketone, acetone, xylene, N-methylpyrrolidone, methyl ethyl ketone, methyl propyl ketone, methyl amyl ketone, and solvent naphtha.

Step b)

In step b) the poly(meth)acrylate containing hydroxy-functional side chains is transesterified or esterified with at least one (meth)acrylate or (meth)acrylic acid or a stabilizer in the presence of an enzyme which
5 catalyzes the transesterification or esterification. Preference is given to carrying out a transesterification with methyl, ethyl, 2-ethylhexyl or butyl (meth)acrylate.

10 The enzymatic transesterification or esterification with a (meth)acrylate or (meth)acrylic acid takes place in general at low temperatures, preferably 10 to 100°C, more preferably 20 to 80°C. The reaction conditions during the enzymatic transesterification or
15 esterification are mild. The low temperatures and other mild conditions prevent the formation of by-products in step b), which may otherwise originate, for example, from chemical catalysts or as a result of unwanted free-radical polymerization of the (meth)acrylate used
20 or of the (meth)acrylic acid used, which can otherwise be prevented only by adding stabilizers.

For the enzymatic reaction (step b)) the product from step a) can be used in general without further
25 pretreatment. If required, the product may be freed from volatiles (e.g., solvents) or additional substances (e.g., solvents) may be added. Specifically, it should as far as possible be free from free-radical initiators or have a low free-radical initiator
30 content.

Preferred enzymes used are hydrolases, especially hydrolases selected from the group consisting of lipases, esterases, and proteases. The enzymes can be
35 used in free form or in immobilized form on a support to which they have been chemically or physically bound. The amount of the enzyme catalyst is preferably 0.1 to 20% by weight, more preferably 1 to 10% by weight,

based on the poly(meth)acrylate containing hydroxy-functional side chains that is used.

5 The reaction time depends among other things on the amount used and on the activity of the enzyme catalyst and the desired degree of conversion, and also on the hydroxy-functional side chain of the poly(meth)-acrylate.

10 The (meth)acrylate used for the transesterification or the (meth)acrylic acid used for the esterification is generally employed in equimolar amounts or in excess in relation to the number of hydroxy-functional side chains in the poly(meth)acrylate. Preference is given
15 to using a molar ratio of (meth)acrylate or (meth)acrylic acid to hydroxy groups in the side chains of the poly(meth)acrylate of 1:1 to 10:2. Higher excesses are not disruptive.

20 Generally speaking, in step b), 20-100%, preferably 40 to 100%, more preferably 60 to 100% of all hydroxy-functional side chains originally present in the poly(meth)acrylate are reacted with a (meth)acrylate or (meth)acrylic acid.

25 Suitable stabilizers, used where appropriate, are selected from the group consisting of 2,6-dibutylphenols such as di-tert-butylphenol, p-cresol, hydroquinone, dimethylhydroquinone, phenothiazines, and
30 phosphorous esters. It is, however, also possible to carry out step b) without using stabilizers.

The reaction can be carried out in all reactors suitable for such a reaction. Reactors of this kind are
35 known to the skilled worker. The reaction takes place with preference in a stirred tank reactor, a fixed bed reactor or a Taylor reactor.

The alcohol formed or the water of reaction formed during the transesterification or esterification can be removed by methods known to the skilled worker: for example, by absorption (with molecular sieve, for
5 example), distillation or pervaporation.

The reaction is continued until the desired conversion, generally 5 to 100%, has been reached. In the case of a reaction regime with simultaneous removal of the water
10 or alcohol formed during the reaction it is possible to achieve higher conversions in shorter reaction times owing to the shifting of the reaction equilibrium.

Following the reaction the enzyme catalyst can be
15 separated off by appropriate measures, filtration or decanting for example, and can if desired be used a number of times.

A further subject of the present specification are UV-
20 curable and/or dual-cure poly(meth)acrylates preparable by the process of the invention. Owing to the mild reaction conditions in the process of the invention it is possible to obtain new kinds of (meth)acryloyl-functional poly(meth)acrylates without the risk of
25 cleavage of the ester bonds in the poly(meth)acrylates by acid catalysis or high temperatures.

These (meth)acryloyl-functional poly(meth)acrylates of the invention are suitable as binders in radiation-
30 curable or dual-cure coating materials: for example, in topcoats such as transparent clearcoat materials, but also in undercoat materials, primers, and surfacers. The (meth)acryloyl-functional poly(meth)acrylates have excellent weather stability. In conjunction with a
35 curing technology (radiation cure or dual cure) it is possible to obtain further advantages: for example, an increase in the scratch resistance of a coating. Particularly decisive, however, is the improvement in

application through use of the (meth)acryloyl-functional poly(meth)acrylates of the invention, since they allow rapid drying.

5 A further subject of the present specification is therefore the use of the (meth)acryloyl-functional poly(meth)acrylates of the invention or of those prepared by the process of the invention as binders in radiation-curable or dual-cure coating materials,
10 preferably in topcoats, more preferably in transparent clearcoat materials.

By "dual-cure" is meant that the materials are curable thermally and with actinic radiation. In the context of
15 the present invention actinic radiation means electromagnetic radiation such as visible light, UV radiation or X-rays, especially UV radiation, and corpuscular radiation such as electron beams.

20 Radiation-curable binders are those curable by means of actinic radiation as defined above, in particular by means of UV radiation.

A further subject of the present specification are
25 coating formulations comprising the (meth)acryloyl-functional poly(meth)acrylates of the invention or those preparable by the process of the invention. The (meth)acryloyl-functional poly(meth)acrylates or the stabilizer-functionalized poly(meth)acrylates can be
30 used both in basecoat materials and in topcoat materials. In view of their particular properties such as the enhancement of scratch resistance in conjunction with high UV stability of a coating their use in topcoats is preferred.

35

Generally speaking, the composition of the topcoat is selected such that the cured topcoat material has a storage modulus E' in the rubber-elastic range of at

least $10^{7.6}$ Pa, preferably of at least $10^{8.0}$ Pa, more preferably of at least $10^{8.3}$ Pa, and a loss factor at 20°C of not more than 1.10, preferably not more than 0.06, the storage modulus E' and the loss factor $\tan\delta$ having been measured by dynamic-mechanical thermoanalysis on homogeneous free films with a thickness of $40 \pm 10 \mu\text{m}$. The loss factor $\tan\delta$ is defined as the quotient of the loss modulus E'' and the storage modulus E' .

Dynamic-mechanical thermoanalysis is a general measurement method for determining the viscoelastic properties of coatings and is described, for example, in Murayama T., Dynamic Mechanical Analysis of Polymeric Material, Elsevier, New York, 1978 and Loren W. Hill, Journal of Coatings Technology, Vol. 64, No. 808, May 1992, pp. 31 to 33. The measurements can be carried out, for example, using the instruments II, MKIII, or MKIV from the company Rheometrics Scientific.

The radiation-curable or dual-cure topcoats preferably have a viscosity at 23°C of < than 100 s efflux time in the DIN4 cup, more preferably < 80 s efflux time in the DIN4 cup. For casting application and roller application the viscosity may also be above this.

The topcoats of the invention comprise in addition to the (meth)acryloyl-functional poly(meth)acrylates of the invention, if desired, one or more photoinitiators and, if desired, customary auxiliaries and additives. Suitable photoinitiators are customary photoinitiators used in radiation-curable or dual-cure coating materials, examples being benzophenones, benzoin ethers, preferably hydroxyacrylic ketones and bis(acyl)phosphine oxides. It is also possible, for example, to use those in commerce under the name Irgacure® 184, Irgacure® 1800, and Irgacure® 500 from

Ciba Geigy, Genocure® MBF from Rahn, and Lucirin® CPO from BASF AG.

Suitable further auxiliaries and additives are, for
5 example, light stabilizers (for example HALS compounds, benzotriazoles, oxalanilith, et cetera), slip additives, polymerization inhibitors, flatting agents, defoamers, leveling agents, and film-forming auxiliaries, cellulose derivatives for example, et
10 cetera. Additionally it is possible to use rheology control components, such as organic urea compounds, urethane urea compounds and/or SiO₂.

The topcoats of the invention are employed in
15 particular as clearcoat materials, so that they normally contain no hiding pigments and no fillers, or only transparent fillers. Also possible, however, is their use in the form of pigmented topcoats. In that case the topcoats additionally comprise pigments.
20 Furthermore, in this case the topcoats may comprise one or more fillers.

A further subject of the present application are therefore topcoats comprising

25 5 to 80% by weight, preferably 10 to 60% by weight, more preferably 20 to 50% by weight of at least one (meth)acryoyl-functional poly(meth)acrylate of the invention or one prepared by the process of the invention,

30

0.5 to 15% by weight, preferably 1 to 10% by weight, more preferably 1 to 5% by weight of at least one photoinitiator,

35 0.5 to 8% by weight, preferably 1 to 6% by weight, more preferably 1 to 4% by weight of further auxiliaries and additives,

0 to 40% by weight, preferably 0 to 30% by weight, more preferably 0 to 25% by weight of pigments,

and 0 to 40% by weight, preferably 0 to 30% by weight,
5 more preferably 0 to 25% by weight of at least one filler, such as transparent metal oxides, BaSO_4 , and waxes.

Preferred (meth)acryloyl-functional poly(meth)acryl-
10 ates, photoinitiators, auxiliaries and additives, and fillers and pigments have already been specified above.

The topcoats of the invention are prepared by mixing the individual components in accordance with methods
15 known to the skilled worker in apparatus known according to the skilled worker.

A further subject of the present specification is therefore a process for preparing the topcoat of the invention, in which the (meth)acryloyl-functional
20 poly(meth)acrylate, the photoinitiator, if desired further auxiliaries and additives, and, if desired, fillers and pigments are mixed with one another.

The topcoats of the invention are generally applied to
25 substrates coated with a basecoat material. They may be applied by what is known as coil coating or by injection molding to the substrates. Such substrates are, for example, metal sheets, or metal strips and plastics of any kind, e.g., automobile bodies and
30 motorcycle parts.

After the topcoat has been applied it is subjected to a radiation cure or dual cure. The equipment and conditions for these curing methods are known from the
35 literature and require no further description (for radiation curing see, for example, R. Holmers, UV and E.B. Curing Formulations for Printing Inks, Coatings

and Paints, SITA Technology, Academic Press, London, United Kingdom 1984).

5 The examples which follow provide further illustration of the invention.

Examples

1. Preparation of a hydroxy-functional binder

10

Formula:

Initial charge	Methyl isobutyl ketone		540.0 g
Monomers	Styrene	10.00% by wt. ¹⁾	123.6 g
	EHA (2-ethylhexyl acrylate)	46.50% by wt. ¹⁾	574.4 g
	HEMA (hexylethyl methacrylate)	27.00% by wt. ¹⁾	336.6 g
	HBA (hydroxybutyl acrylate)	15.00% by wt. ¹⁾	185.2 g
	AA (acrylic acid)	1.50% by wt. ¹⁾	18.6 g
rinse	Methyl isobutyl ketone		5.0 g
Initiator	tert-Butyl peroxy-2-ethylhexanoate	8 % by wt. ²⁾	98.8 g
	Methyl isobutyl ketone		74.2 g
rinse	Methyl isobutyl ketone		46.6 g
End			2000.0 g

15 1) based on the sum of the components styrene, EHA, HEMA, HBA, AA

2 based on the sum of the components styrene, EHA, HEMA, HBA, AA

Procedure:

20 Weigh out initial charge and heat to 110°C. At constant temperature, meter monomers and initiator into the reactor at a uniform rate. After 4 hours the monomer

feed is at an end. After 4.5 hours the initiator feed is at an end. After the end of the metering of initiator polymerization is continued for 1 hour, followed by cooling and discharge of the reaction mixture obtained.

*end values:

	Solids (1 h, 130°C):	66.3%
10	OH number (theoretical) total:	174.8 mg/g (determined to DIN 43402)
	OH number (practical):	165 mg/g (determined to DIN 53246)
	GC (residual monomer content) ³⁾ :	EHA 0.3%; AA <0.3% all others < 0.1%
15	GPC ⁴⁾ Mn ⁵⁾	5829
	M _w ⁶⁾	20 722
	M _w /M _n ⁷⁾	3.55

- 3) GC = gas chromatography
- 20 4) GPC = gel permeation chromatography (with polystyrene standard)
- 5) M_n = number-average molecular weight
- 6) M_w = weight-average molecular weight
- 7) M_w/M_n = polydispersity

25

2. Preparation of a UV-active polyacrylate

Batch: 300 ml of polymer solution in methyl isobutyl ketone from Example 1

30 300 g of methyl acrylate (MA)

150 mg of methoxyphenol

150 g of 5 Å mole sieve

30 g of Novozym® 435 (immobilized lipase from Candida antarctica from the company Novozymes)

35

The components stated are stirred at 40°C for 72 hours. The reaction mixture is subsequently filtered and the polyacrylate obtained is washed with methyl isobutyl

ketone (MIK). The excess MA and MIK is removed in vacuo on a rotary evaporator at 60°C to 70°C. This gave 227 g of target product. The fraction of the acrylated hydroxy groups was determined as being about 34% by means of the OH number.

The OH number was determined in accordance with a method which is known in the prior art (DIN 53240, Part 2).

3. Preparation of a UV coating formulation

a) Stock varnish

inventive UV polyacrylate	32.5
Sartomer® 399	30.6
Thixharz® SCA	11.5
(basis: benzylamine/hexamethylene diisocyanate)	
Irgacure® 184 (photoinitiator)	0.8
Lucirin® TPO (photoinitiator)	0.4
Byk® 358 (leveling assistant)	0.2
Tinuvin® 292 (free-radical scavenger)	1.0
Tinuvin® 400 (UV absorber)	1.0
Butyl acetate	22.0

b) Curing agent mixture

A curing agent mixture composed of 72.7 parts of Roskydal® UA VP LS 2337 (unsaturated isophorone diisocyanate), 18.2 parts of Roskydal® UA VP FWO 3003 77 and 9.1 parts of butyl acetate is added (the parts are parts by weight).

The components were mixed with a dissolver.

4. The coating formulation was applied by spray application.

5. Variation of the hydroxy-functional units

As the following examples show, different hydroxy-functional units can be used. The examples mentioned are not, however, intended to be any restriction. The polymer solutions were prepared by methods
5 corresponding to Example 1.

10 g of polymer solution, 10 g of methyl acrylate, 5 g of mole sieve (5 Å) and 1 g of immobilized lipase (Novozym® 435) were shaken at 40°C for 72 hours. After
10 filtration and concentration, the conversion was determined by way of the OH number.

Polymer solution	Esterified unit	Conversion [%]
2	Hydroxyethyl acrylate	34
3	Hydroxyethyl acrylate	41
4	Hydroxyethyl methacrylate	12
5	Hydroxyethyl methacrylate	22
6	Hydroxyethyl methacrylate	47
7	Hydroxybutyl acrylate	67
8	Hydroxybutyl acrylate	80

6. Reaction optimization

15

It was possible to optimize the reaction conditions by varying the reaction time, the added amount of methyl acrylate, and mole sieve. The polymer solutions shown in the table under Example 5 were reacted under the
20 following optimized reaction conditions.

10 g of polymer solution, 2 g of methacrylate, 2 g of mole sieve and 1 g of Novozym® 435 were shaken at 40°C for 24 hours. After filtration and concentration, the
25 conversion was determined by way of the OH number.

Polymer solution	Conversion [%]
2	23
3	39

5	6
6	17
7	40
8	46

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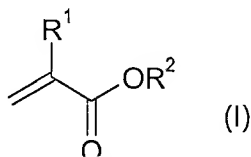
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Claims

5 1. A process for preparing poly(meth)acrylates curable with actinic radiation and/or dual-cure poly(meth)acrylates, comprising the following steps:

10 a) preparing a poly(meth)acrylate containing hydroxy-functional side chains by polymerizing

aa) at least one (meth)acrylate of the general formula (I) as component A

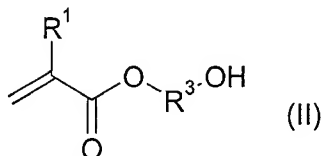


15 in which

R^1 is H, CH_3 or CH_2OH and

20 R^2 is an alkyl radical which is unsubstituted or substituted by functional groups such as acrylic, ether, amino, epoxy, halogen or sulfonic acid groups, and

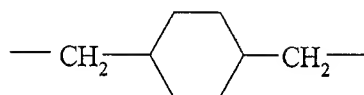
25 ab) at least one hydroxyalkyl (meth)acrylate of the general formula (II) as component B



in which

R^1 is H, CH_3 or CH_2OH and

R^3 is $-(CH_2)_n-$, $-CH_2-CH(CH_3)-CH_2-$ or
 $-CH_2CH(CH_3)-$ or $-CH(CH_3)CH_2-$ or



n is at least 2, and

5

ac) if desired, further comonomers,
copolymerizable with the (meth)acrylates
of the general formula (I) and (II), as
component C, and

10

ad) if desired, auxiliary monomers as
component D;

and

15

b) transesterifying or esterifying the
poly(meth)acrylate containing hydroxy-
functional side chains with a (meth)acrylate
or (meth)acrylic acid in the presence of an
enzyme which catalyzes the trans-
esterification or esterification.

20

2. A process as claimed in claim 1, wherein step a)
is carried out using

25

- 10 to 80% by weight of component A,
- 10 to 80% by weight of component B,
- 0 to 50% by weight of component C, and
- 0 to 15% by weight of component D.

30

3. A process as claimed in claim 1 or 2, wherein
enzymes used in step b) are hydrolases selected
from the group consisting of lipases, esterases,
and proteases.

4. A process as claimed in any of claims 1 to 3, wherein step b) is carried out using methyl, ethyl, 2-ethylhexyl or butyl (meth)acrylate.
- 5 5. A process as claimed in any of claims 1 to 4, wherein the temperature at which step b) is conducted is 20 to 100°C, preferably 20 to 80°C.
- 10 6. A process as claimed in any of claims 1 to 5, wherein component B is selected from the group consisting of 2-hydroxyethyl (meth)acrylate, 2-hydroxypropyl (meth)acrylate, and hydroxybutyl (meth)acrylate.
- 15 7. A process as claimed in any of claims 1 to 6, wherein 5 to 100% of the in the side chain of the poly(meth)acrylate prepared in accordance with step a) have been (meth)acrylated.
- 20 8. Poly(meth)acrylates curable with actinic radiation and/or dual-cure poly(meth)acrylates preparable by a process as claimed in any of claims 1 to 7.
- 25 9. The use of poly(meth)acrylates curable with actinic radiation and/or dual-cure poly(meth)-acrylates as claimed in claim 8 or prepared by a process as claimed in any of claims 1 to 7 as a component in the preparation of dispersions or as a component in coating formulations, preferably in
30 actinic-radiation-curable and/or dual-cure coatings or topcoats, more preferably in transparent clearcoat materials.
- 35 10. A topcoat containing
 - 5 to 80% by weight of at least one poly(meth)acrylate curable with actinic radiation and/or dual-cure poly(meth)acrylate

- as claimed in claim 8 or prepared as claimed
in any of claims 1 to 7,
- 0.5 to 15% by weight of at least one
photoinitiator,
- 5 0.5 to 8% by weight of further auxiliaries
and additives,
0 to 40% by weight of pigments, and
0 to 40% by weight of at least one filler.
- 10 11. A process for preparing a coating formulation as
claimed in claim 10, in which the individual
components are mixed with one another.
12. The use of a coating formulation as claimed in
15 claim 10 as a topcoat.

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Abstract

The present invention relates to processes for preparing actinic-radiation-curable and/or dual-cure poly(meth)acrylates by preparing a poly(meth)acrylate containing hydroxy-functional side chains and trans-esterifying or esterifying the poly(meth)acrylate containing hydroxy-functional side chains with a (meth)acrylate or (meth)acrylic acid. The present invention further relates to the actinic-radiation-curable and/or dual-cure poly(meth)acrylates themselves and to the use of the actinic-radiation-curable and/or dual-cure poly(meth)acrylates in the preparation of dispersions or as a component in coating formulations and topcoats comprising at least one actinic-radiation-curable and/or dual-cure poly(meth)acrylate.